WEST Search History

DATE: Thursday, July 17, 2003

Set Name side by side	Query	Hit Count	Set Name result set
DB = USPT, PG	SPB,JPAB,EPAB; PLUR=YES; OP=ADJ	7	
L13	L12 and 13	4	L13
L12	rip14 or rip-14	. 8	L12
L11	rip[14 or rip-14	2	L11
L10	L9 and l3	18	L10
L9	bile acid receptor	28	L9
L8	L6 and 13	1	L8
L7	L6 and 11	0	L7
L6	nr1h4	13	L6
L5	L4 and 13	29	L5
L4	fxr	155	L4
L3	antisense or anti-sense	32368	L3
L2	L1	19468	L2
DB=USPT; PLUR=YES; OP=ADJ			
L1	antisense or anti-sense	19468	L1

END OF SEARCH HISTORY

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                 PCTFULL now contains images
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        Mar 04
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                 PATDPAFULL now available on STN
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        Mar 24
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                 structures available in REGISTRY
NEWS 10
                 Display formats in DGENE enhanced
        Apr 11
NEWS 11
        Apr 14
                 MEDLINE Reload
NEWS 12
        Apr 17
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NEWS 14
                 New current-awareness alert (SDI) frequency in
                 WPIDS/WPINDEX/WPIX
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        Apr 28
                 RDISCLOSURE now available on STN
NEWS 16
        May 05
                 Pharmacokinetic information and systematic chemical names
                 added to PHAR
NEWS 17
                 MEDLINE file segment of TOXCENTER reloaded
        May 15
NEWS 18
        May 15
                 Supporter information for ENCOMPPAT and ENCOMPLIT updated
NEWS 19
        May 19
                 Simultaneous left and right truncation added to WSCA
NEWS 20
        May 19
                 RAPRA enhanced with new search field, simultaneous left and
                 right truncation
NEWS 21
        Jun 06
                 Simultaneous left and right truncation added to CBNB
        Jun 06
NEWS 22
                 PASCAL enhanced with additional data
NEWS 23
        Jun 20
                 2003 edition of the FSTA Thesaurus is now available
NEWS 24
        Jun 25
                HSDB has been reloaded
        Jul 16 Data from 1960-1976 added to RDISCLOSURE
NEWS 25
             April 4 CURRENT WINDOWS VERSION IS V6.01a, CURRENT
NEWS EXPRESS
              MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
              AND CURRENT DISCOVER FILE IS DATED 01 APRIL 2003
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=> file medline caplus biotechno biosis scisearch embase

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION

0.42

0.42

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=> s fxr

L2 878 FXR

FULL ESTIMATED COST

=> s nr1h14

L3 0 NR1H14

=> s nrih4

L4 0 NRIH4

=> s nr1h4

L5 70 NR1H4

=> s rip14

L6 17 RIP14

=> s rip-14

L7 3 RIP-14

=> s bile acid receptor

L9 177 BILE ACID RECEPTOR

=> s 12 or 15 or 16 or 17 or 18 or 19 L10 983 L2 OR L5 OR L6 OR L7 OR L8 OR L9

=> s 110 and 11

L11 3 L10 AND L1

=> dup rem 111

PROCESSING COMPLETED FOR L11

L12 3 DUP REM L11 (0 DUPLICATES REMOVED)

- L12 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2003 ACS
- AB Antisense compds., compns. and methods are provided for modulating the expression of human FXR (farnesoid X receptor). The compns. comprise antisense compds., particularly antisense oligonucleotides, targeted to nucleic acids encoding human FXR. Methods of using these compds. for modulation of human FXR expression and for treatment of cardiovascular disease and atherosclerosis assocd. with expression of human FXR are provided.
- L12 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2003 ACS
- The present invention relates to a human and mouse novel nuclear receptor called "L66" or also FXR-.beta. (farnesoid X receptor .beta.), a homolog of the FXR-.alpha. which is a prototypical type 2 nuclear receptor. The invention also relates to the isolated nucleic acid sequence of L66 and the isolated protein thereof. The invention further relates to processes for isolating and/or producing the nucleic acid or the protein as well as methods of use of the receptor L66. The invention also provides the sequence of mouse gene L66 and reverse complement sequence. The invention also relates to screening drugs which are capable of inhibiting the cellular function of the nuclear receptor L66 in cells, including antibody, RNA, antisense oligo and ribozyme. The invention also relates to expression of L66 in tissues and cell lines.
- L12 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2003 ACS
- Fibrosis is a consequence of injury characterized by accumulation of excess collagen and other extracellular matrix components, resulting in the destruction of normal tissue architecture and loss of function. Spl was originally described as a ubiquitous transcription factor. It is involved in the basal expression of extracellular matrix genes and may, therefore, be important in fibrotic processes. To evaluate the effect of Sp1 blockade on the expression of extracellular matrix genes, clones of NIH 3T3 fibroblasts stably transfected with an antisense Sp1 expression vector. Simultaneously reduced expression of
 several extracellular matrix genes as compared with mock-transfected clones was noted using differential hybridization of cDNA microarrays, without significant alteration in cell growth. Transfection of human dermal fibroblasts with several extracellular matrix gene (COL1A1, COL1A2, COL3A1, COL5A2, COL7A1, TIMP-1, and decorin) promoter/reporter constructs demonstrated that anti-sense Spl-induced redn. of extracellular matrix gene mRNA steady-state levels results from transcriptional repression, consistent with the role of Sp1 as a transcription factor. Decoy Sp1 binding oligonucleotides inhibited COL1A2 promoter activity both in cultured fibroblasts and in vivo, in the skin of transgenic mice, which have integrated a mouse COL1A2 promoter/luciferase reporter gene construct. These results indicate that targeting Sp1 efficiently blocks extracellular matrix gene expression, and suggest that such an approach may represent an interesting therapeutic alternative toward the treatment of fibrotic disorders.
- => s ribozyme or ribozymes L13 23191 RIBOZYME OR RIBOZYMES
- => s 110 and 113 L14 1 L10 AND L13
- => 114 not 112 L14 IS NOT A RECOGNIZED COMMAND The previous command name entered was not recognized by the system. For a list of commands available to you in the current file, enter

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"HELP COMMANDS" at an arrow prompt (=>).
=> s 114 not 112
L15
              0 L14 NOT L12
=> d 1-3 l12
L12 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2003 ACS
     2003:417857 CAPLUS
AN
     139:17574
DN
     Use of antisense oligonucleotides to gene encoding human
     farnesoid X receptor for treatment of cardiovascular disease and
     atherosclerosis
     Monia, Brett P.; Watt, Andrew T.
IN
     Isis Pharmaceuticals, Inc., USA
PA
     PCT Int. Appl., 127 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 1
     PATENT NO.
                       KIND DATE
                                               APPLICATION NO. DATE
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PΙ
     WO 2003044167
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                                               WO 2002-US36691 20021113
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     US 2003109467
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PRAI US 2001-2491
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     ANSWER 2 OF 3 CAPLUS COPYRIGHT 2003 ACS
L12
     2002:220785 CAPLUS
AN
     136:274299
DN
TI
     Human and mouse nuclear receptor L66, protein and cDNA sequences and
     recombinant production
     Casari, Georg; Hoefer, Michael; Jackson, David; Kranz, Harald; Otte,
IN
     Kerstin; Remmel, Bettina; Suckow, Joerg
PA
     Lion Bioscience A.-G., Germany
     PCT Int. Appl., 136 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
     PATENT NO.
                       KIND DATE
                                               APPLICATION NO. DATE
                       _ _ _ _
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     WO 2002022817
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AU 2002013893

A5 20020326

AU 2002-13893 20010907

EP 1317542 A2 20030611 EP 2001-982261 20010907
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

PRAI EP 2000-120370 A 20000916 EP 2001-111658 A 20010514 WO 2001-EP10323 W 20010907

- L12 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2003 ACS
- AN 2001:402491 CAPLUS
- DN 135:236379
- TI Blocking Sp1 transcription factor broadly inhibits extracellular matrix gene expression in vitro and in vivo: implications for the treatment of tissue fibrosis
- AU Verrecchia, Franck; Rossert, Jerome; Mauviel, Alain
- CS INSERM U532, Hopital Saint-Louis, Paris, 75475, Fr.
- SO Journal of Investigative Dermatology (2001), 116(5), 755-763 CODEN: JIDEAE; ISSN: 0022-202X
- PB Blackwell Science, Inc.
- DT Journal
- LA English
- RE.CNT 61 THERE ARE 61 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT